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**Bridge, Jennifer A.; Overgaard, Nana Haahr; Steptoe, Raymond; Frazer, Ian H.; Wells, James W.**

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# **Altering the balance between immune activation versus regulation in the skin to promote CD8<sup>+</sup> T-cell activity within epithelial cancers**

Jennifer A. BRIDGE<sup>1</sup>, Nana H. OVERGAARD<sup>1</sup>, Raymond J. STEPTOE<sup>1</sup>, Ian H. FRAZER<sup>1</sup> and James W. WELLS<sup>1</sup>.

<sup>1</sup>The University of Queensland Diamantina Institute, The University of Queensland, Translational Research Institute, Brisbane, QLD 4102, AUSTRALIA, Email: [j.wells3@uq.edu.au](mailto:j.wells3@uq.edu.au)

The Human Papilloma Virus (HPV) 16 is a high-risk HPV known to be a causative agent in numerous cancers including cervical cancer. While prophylactic vaccines exist to combat the spread of HPV16, successful therapeutic vaccines to combat established HPV16-associated disease remain elusive. The expression, in a mouse model ("E7"), of the HPV16 E7 gene in keratinocytes under the control of the K14 promoter, leads to a local immune suppressive environment, as evidenced by the lack of graft rejection when E7 skin grafts are placed on WT recipient mice. Furthermore, well healed (>30 days) E7 skin grafts are not rejected when mice are immunised with E7 peptide in combination with Quil A- or CASAC-based adjuvants. This is despite a substantial increase in E7 peptide/H-2D<sup>b</sup> pentamer staining in the blood, and marked killing of E7-peptide expressing TC-1 cells when injected i.v., confirming that CD8 T-cells respond to vaccination and differentiate into CTL capable of killing E7-expressing target cells. We hypothesised that the removal of regulatory T-cells (T-reg) might lead to E7 graft rejection in immunised mice. The co-administration of an anti-CD4-depleting antibody at the time of immunisation led to rejection of ~50% of grafts. To confirm a role for T-reg, E7-grafted T-reg-deficient Rag1<sup>-/-</sup> mice received purified donor CD8 T-cells from E7-vaccinated WT mice. FACS staining of Rag1<sup>-/-</sup> lymph nodes 30 days post CD8<sup>+</sup> T-cell transfer confirmed the absence of classical CD4<sup>+</sup>FoxP3<sup>+</sup> Treg, however the E7 grafts did not reject. As in the WT mice however, rejection could be induced through the co-administration of an anti-CD4 antibody. The data suggest that the removal of a CD4<sup>+</sup>, non T-reg cell, leads to CD8<sup>+</sup> T-cell activity in the skin as evidenced by E7 skin graft destruction.